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(54) Surgical devices for imposing a negative pressure to stabilize cardiac tissue during surgery

(57) The invention is devices and techniques which use a negative (suction) pressure or vacuum, applied through a surgical instrument, to fix the position of a portion of the surface of a beating heart so that a surgical procedure can be more easily performed. The devices apply a negative pressure at several points on the outer surface of the heart such that a portion of the heart is fixed in place by the suction imposed through the surgical instrument. Because the instrument fixes the position of the tissue, and because the instruments remain at a constant distance from the particular portion of the heart where the surgery is being performed, the device may also serve as a support or platform so that other surgical instruments or devices can be advantageously used at the site. In certain preferred embodiments, the devices described herein have structures to facilitate the use of additional surgical instruments such that the placement of negative pressure device permit the surgeon to advantageously manipulate the other instruments during surgery. The negative pressure is preferably imposed through a plurality of ports which may be disposed in a substantially planar surface of the instrument which contacts the cardiac tissue.

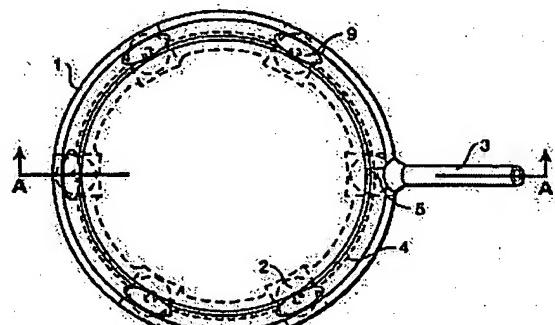


FIG. 1

Description

Surgeries to treat disease in the heart, particularly blockages in coronary vessels, are becoming increasingly common and necessary to treat atherosclerosis and other conditions causing reduced blood flow to the muscles of the heart. For many years, surgeons have performed "open heart" surgery to repair defects in the muscles of the heart and the associated cardiovascular system. As these procedures have become more common and more costly, a need has developed for techniques to make cardiac surgical procedures less traumatic to the patient. The necessity of attaching a surgical patient to a cardiopulmonary bypass (CPB) apparatus is a primary contributor to the trauma inherent in traditional procedures. To attempt to alleviate the trauma and side effects of CPB, surgeons have begun performing cardiac surgeries without stopping the heart. To successfully perform such surgery, several challenges must be met. One particular problem confronting the surgeon who operates on the beating heart is the difficulty in performing extremely delicate surgical procedures while the contractions of the heart muscles cause the surface of the heart to continuously move.

To attempt to restrict the motion of heart at the particular area where the surgeon is working, the surgeon may pass at least a pair of sutures through the exterior tissue layers of the heart. By pulling the sutures in opposite directions, the tissue is stretched, and the motion caused by the contractions of the heart muscles is reduced or partially compensated. This technique is not completely effective in restricting the natural motion of the heart and requires extra time to place the sutures, and, additionally, may cause damage to the cardiac tissue when the sutures are placed or manipulated. Preferably, the surgeon would be able to fix the motion of the cardiac tissue containing or adjacent to an area where surgery is to be performed without the need to attach or manipulate additional sutures. The ability to fix the position of the cardiac tissue in a selected region of the heart would permit the surgeon to perform delicate surgical procedures on the beating heart while the portion of the heart on which the surgery is performed remains substantially motionless throughout the procedure.

SUMMARY OF THE INVENTION

This invention is devices and techniques which use a negative pressure (vacuum) applied through a surgical instrument, to fix the position of a portion of the surface of a beating heart so that a surgical procedure can be more easily performed. The devices disclosed herein apply a negative pressure at several points on the outer surface of the heart such that a portion of the exterior tissue of the heart is fixed in place by the suction imposed through the surgical instrument. Because the negative pressure introduced through the instrument

tion of the heart where the surgery is being performed. In this configuration, the device may also serve as a support or platform so that other surgical instruments or devices can be advantageously used at the site. In certain preferred embodiments, the devices described herein have structures to facilitate the use of additional surgical instruments such that the placement of the negative pressure instrument permits the surgeon to advantageously manipulate the other instruments used during the surgery.

The negative pressure is preferably imposed through a plurality of ports which may be disposed in a substantially planar surface of the instrument which contacts the cardiac tissue. The ports are preferably oriented such that the pressure is applied at several points over the target area to fix the position of the tissue and to reduce any trauma to the tissue caused by the negative pressure.

DESCRIPTION OF THE FIGURES

Figure 1 is an embodiment of the invention having a substantially annular housing with a plurality of suction ports disposed about the periphery of the instrument and having openings in the bottom surface which contacts the heart.

Figure 2 is a dome-shaped or semi-spherical embodiment having a plurality of suction ports disposed about the periphery of the bottom surface and having several instrument ports in the dome portion through which additional surgical instruments may be introduced, positioned, or manipulated.

Figure 3 is a section of a substantially circular embodiment showing a preferred configuration for the suction ports and a pressure conducting chamber for introducing the negative pressure to each suction port.

Figure 4 is an embodiment of the instrument, in use, which is fixed on the surface of the heart and has additional surgical instruments operably associated therewith to facilitate a graft or a source artery being attached to a target vessel form an anastomosis, for example, between the internal mammary artery (IMA) and the left anterior descending (LAD) artery.

Figure 5 is an embodiment of the invention wherein the suction ports for imposing the negative pressure are affixed to a shaft, which may be part of a hand-held device, and are contained in a block where a plurality of individual suction ports are arranged in an array.

Figure 6 is a plurality of suction ports contained within a block having vacuum lines attached thereto for introducing a negative pressure to each suction port.

Figures 7 and 7a are sectional views of a block showing a configuration for the suction ports and the pressure conducting space.

Figure 8 is an array of suction ports having vacuum tubes which run the length of the block and are oriented to be substantially perpendicular to a passage space leading to each suction port.

portion thereof, having a removable block, wherein the instrument has a receiving means to reversibly receive the block.

DETAILED DESCRIPTION OF THE INVENTION

This invention is surgical instruments and techniques which advantageously apply a negative pressure to the surface of the heart so that a portion thereof is maintained at a fixed position during a surgical procedure. The negative pressure is introduced to the instrument and is applied at several points over the surface of the heart proximate to or surrounding the portion of the heart whose position is desired to be fixed during the procedure. The instruments feature several suction ports which are brought into contact with the heart, followed by the application of a negative pressure through the instrument, to fix the position of a region of cardiac tissue based on the placement of the instrument. The instruments may also contain a sealed, airtight, pressure conducting chamber operably connected to a pressure inlet for communicating the negative pressure to the suction ports. Alternatively, each suction port may have a dedicated vacuum line attached thereto.

The shape of the instrument may be varied depending on the particular application or the clinical diagnosis for an individual patient. In some embodiments, the shape of the instrument is defined by a housing forming a complete or partial, substantially annular, ring-shaped housing having the suction ports disposed about the periphery of the bottom surface of the housing. The suction ports are contained within the base of the instrument and the opening of the suction ports are contained in the bottom surface of the instrument which may be substantially planar or may be shaped to conform to the surface of the heart.

In another embodiment, the operative portion of the instrument may be defined by one or more arrays of suction ports which are substantially linear. The suction ports may be contained in a block which has at least one vacuum line attached thereto. This design is particularly suitable for an instrument having a shaft affixed thereto for positioning the block containing the suction ports. The shaft may be fixed to a rigid support during the procedure or may be part of a hand-held instrument having a handle structure adapted to be grasped by the human hand. In a preferred embodiment, the hand-held instrument contains a pair of shafts having a block and suction port assembly at each end thereof. The shafts are connected at an intermediate portion by a pivot which allows the suction port assemblies to move relative to one another, to be oriented and manipulated by hand, and to be locked into place in a desired configuration.

An embodiment having more than one movable member in which suction ports are contained offers the advantage that a negative pressure may be first imposed through the suction ports of each movable

individual members which causes the tissue to be stretched or oriented such that one portion of the cardiac tissue is fixed in position by one movable member and can be oriented relative to another portion fixed by a second movable member.

The negative or vacuum pressure imposed may be varied depending on the design of the instrument, the orientation of the ports, and the amount of pressure needed to hold a particular region of the heart in place. When manipulating the instruments of this invention, it is not desired to exert a downward force on the instrument once the instrument engages the cardiac tissue because the tissue could be damaged by being drawn into the suction ports, thus risking interruption of blood flow and ischemic or reperfusion injury to the cardiac tissue. However, once a negative pressure is imposed, the instrument may be drawn away from the heart such that the portion of the surface tissue fixed by the suction ports is slightly elevated relative to the remainder of the heart.

DETAILED DESCRIPTION OF THE INVENTION

Referring to Figures 1 and 3, an embodiment of the invention is an instrument comprised of an annular housing 1 which could have an alternate shape depending on the design and clinical application of the instrument. For example, the body of the instrument has a housing which may be a portion of a circle, an oval, semi-oval, U-shape, or linear member. The portion of the housing 1 which contains the suction ports 2 has a bottom surface 6 which rests against the surface of the heart and therefore should be substantially planar or curved to the extent necessary to simultaneously bring the suction ports 2 into conforming contact with the heart. Referring to Figures 1 and 3, one embodiment of the invention has suction ports 2 equally spaced about the circumference of the annular housing 1. For purposes of stability, it is preferred that each suction port 2 be substantially equidistant from each adjacent port and their location spread over the entire portion of the instrument at the points of contact to the heart by the bottom portion of the instrument to more stably affix the instrument to the surface of the heart when a negative pressure is imposed. As described in detail below, the housing 1 may also have one or more instrument ports 9 to facilitate introducing a surgical instrument to the site of the surgery to function on or in proximity to the fixed portion of the cardiac tissue.

The interior of the housing 1 may be further comprised of a means for introducing a negative pressure to the suction ports 2. For example, each suction port 2 may have a dedicated vacuum line 3 for introducing a negative pressure to each suction port 2. However, it is preferred that a single vacuum line 3 introduce the negative pressure via an inlet 5 which leads to an airtight, sealed, and pressure conducting chamber 4 contained within the annular housing 1 which in turn communi-

Thus, by connecting a negative pressure source to the inlet 5, the negative pressure is introduced to the instrument through inlet 5, thereby creating a negative pressure in the pressure conducting chamber 4 which is communicated to each suction port 2. The housing may also have at least one instrument port 9 comprised of an opening that preferably traverses the width of the housing and is shaped to receive an instrument. In use, the surgeon may rely on the housing 1 as a platform for other instruments which may advantageously be used at the portion of the heart fixed in place by the negative pressure. The instrument port 9 may be a simple opening in the housing 1 or may be designed to operably receive a specific instrument as described in more detail below.

In use (See Figure 4), the instrument is gently positioned on the surface of the heart by manipulating the position of the housing 1 such that the suction ports 2 rest against the cardiac tissue. Once the instrument is positioned on the surface of the heart, the negative pressure is applied through vacuum line 3 and inlet 5 while the housing may be gently manipulated to insure that the negative pressure is causing the cardiac tissue to become fixed to the suction ports 2. Once the suction ports become functionally attached to the surface of the heart, the portion of the surface of the heart becomes fixed relative to the instrument. Once the negative pressure is applied, the instrument may be attached to a stable support such as a rib retractor or other structure which does not move relative to the beating heart.

Referring to Figure 2, a dome-shaped or semi-spherical embodiment of the invention has a plurality of suction ports 2 spaced about the periphery of the bottom surface 6 of the dome portion 8 such that the entire instrument is fixed to the cardiac tissue at the point of contact by each of the several suction ports 2. As with the above embodiment, it is preferred that each suction port 2 be pneumatically connected via an air-tight pressure conducting chamber 4. The base of the instrument is comprised of a substantially flat bottom surface 6 wherein the opening of each of the suction ports 2 is flush at the bottom surface 6. The bottom surface 6 is preferably substantially flat because the bottom surface 6 will engage the surface of the heart when the negative pressure is imposed. Alternatively, depending on the size of the instrument and the location of placement on the surface of the heart, the bottom surface 6 may be contoured so that the suction ports 2 may engage a curved surface of the heart. The bottom surface 6 may also have a separate contact layer 7 to cushion the contact between the instrument and the heart tissue and to facilitate forming a tight seal when the negative pressure is imposed. The contact layer 7 may cover substantially the entire bottom surface 6 proximate to the openings of the suction ports 2. If the contact layer material surrounds the openings of the suction ports, it is preferable that the material not be air permeable to prevent the negative pressure from passing through the

attached at the periphery of the bottom surface 6. The available materials for the contact layer 7 include the well-known and commercially available medical plastics such as teflon, silicon, and others which are pliable and biocompatible.

The dome-shaped or semi-spherical embodiment of the invention preferably has at least one instrument port 9 which may be placed in any of several locations but which is preferably located in the dome portion 8 of the apparatus. The instrument port 9 facilitates introducing the functional portion of an instrument 10 into the interior of the dome portion 8, such that the instrument 10 can perform any of several functions on or about the cardiac tissue. The instrument 10 could include a cutting apparatus, visual means, such as a scope or light, suturing instruments, suction, blowing, or irrigation apparatus or any like instrument used during a surgical procedure. Multiple instrument ports 9 disposed in the dome portion 8 allow several instruments 10 to be introduced to the surgical site from numerous directions and to be fixed in place relative to the heart.

The instrument ports 9 may be comprised of only a simple opening in the dome portion 8. Alternatively, the instrument ports 9 may also have a flexible or rigid shaft 11 or other attachment means fixed to the dome portion 8 to facilitate introducing an instrument or a member associated therewith such as wires, tubes, cables, etc. which comprise or are used to perform the function of the instrument 10. The shaft 11 may also comprise the inlet (not shown) for introducing negative pressure to the pressure conducting chamber 4. Because the dome portion 8 remains at a fixed distance to the heart, the instrument ports 9 or shaft 11 may have a collar 23 or stop associated therewith such that the distance between the instrument 10 and the heart can be predetermined and fixed by, for example, abutting a stop 12a on the instrument 10 against a stop 12b or collar on the instrument port 9.

The instrument ports may also contain a locking means, which may be magnetic or suction-driven, so that the instrument 10 can be locked into place on the dome portion 8. For example, the surgeon may view the procedure via a scope 22 which communicates an image to a video monitor. This invention may be advantageously used to establish and maintain an optimal position for the scope by inserting the scope 22 through the instrument port and then fixing the position of the scope 22. The end of the scope 22 may have a collar 23 or other stop mechanism near its end, such that the scope 22 may be introduced through the instrument port 9 wherein the movement of the scope 22 toward the heart is terminated by the contact between the collar 23 of the scope 22 and the periphery of the instrument port 9.

Referring to Figure 3, a sectional view through line A-A of Figure 1 shows the interior of the pressure conducting chamber 4 and associated passages or conduits associated with the suction ports 2 such that the

4 is introduced from vacuum line 3, via inlet 5, and ultimately to the suction ports 2 to provide the means for introducing the negative pressure. The suction ports 2 have a substantially circular opening 12 disposed in the bottom surface 6. Although this embodiment has a single inlet 5 such that the pressure is introduced to each suction portion 2 via the pressure conducting chamber 4, one or more of the suction ports 2 may each optionally have an independent inlet 5' for a separate vacuum line 3'.

As noted above, the suction ports 2 are disposed within a bottom surface 6 which is preferably substantially flat. As noted above, the bottom surface 6 of the housing 1 may be a continuous ring or other annular shape which contacts the heart about the entire periphery of housing 1, but may also be comprised of a plurality of individual bases 13 which contain the one or more suction ports 2 and which contact the heart at several independent points which may be co-planar or which may be adapted to the contours of the heart.

Referring to Figure 4, Figure 4 shows an embodiment of the invention in use in a coronary artery bypass graft (CABG) procedure where an anastomosis is formed between the internal mammary artery (IMA) 13 and the left anterior descending artery (LAD) 14 which is held open by vessel retractors 16a and 16b. One end of the anastomosis is sewn to the LAD 14 by sutures 17 being manipulated by instrument 10. A vacuum line 3 is attached to inlet 5, to introduce a negative pressure to the pressure conducting chamber 4. An instrument 10, which in this example is manipulating suture 17 for sewing the anastomosis at the LAD 14, is introduced via instrument port 9a located in the housing 1 of the apparatus. An instrument port 9a has a shaft 18 disposed within the instrument port 9a to facilitate positioning the instrument 10 relative to both the housing 1 and to the surgical site. The shaft 18 traverses all or a portion of the instrument port 9a and may be flexible such that the shaft 10 can be oriented in a fashion to direct the instrument 10 to the desired point within the surgical field. The shaft 18 may also be incorporated into a pivot 24 of any of several configurations, including a ball 25 and socket 26 joint having a passage 27 running axially through the ball 25 wherein the shaft 18 is contained in the passage 27 such that the ball 25 is rotated within the socket 26 to rotate the instrument 10 about the pivot 24 to position the functional end of the instrument 10. An example of another instrument 19 is inserted through instrument port 9b by passing the instrument 19 through a rigid or flexible shaft 12 which is disposed within the instrument port 9b. In this example, the instrument 19 passes through the shaft 12 such that the position of the functional end of the instrument 19 is fixed. In the example shown in Figure 4, the instrument 19 is being used to grasp the IMA 13 proximate to the attachment to the LAD.

Referring to Figure 5, pursuant to this invention, the suction ports 2 may be incorporated into a hand-held

suction port assembly 22 containing the suction ports 2, at least one vacuum line 3, and a block 23 wherein the suction ports 2 are contained. This embodiment is adapted for, may be manipulated by, a surgeon or surgical assistant by hand grips 24 at the proximal ends of the shaft or the hand grips 24 may be replaced by a conventional attachment (not shown) for fixing the shaft 21 to a stable support such as a surgical retractor or other such fixed structure available during the surgical procedure. In a preferred embodiment, first and second shafts 21 are interlinked by a pivot link 25 at an intermediate point in the shaft 21 such that each suction port assembly may be positioned independently of one another either before or after the negative pressure is applied. In this configuration, the cardiac tissue which is functionally attached to the instrument at the points of contact with the suction port assembly 22 may be drawn apart by imposing a negative pressure through the suction ports 2 and then independently manipulating the rigid shafts 21, or proximally disposed hand grips 24 of the instrument 20 such that each suction port assembly 22 is moved a distance from the other. The suction port assembly 22 is comprised of the block 23, in which the suction ports 2 are contained, and a receiving means 29 located at the terminal (lower) or distal end of the shaft 21 which is shaped to receive the block 23 which is in turn adapted to receive a vacuum line 3 to permit the introduction of negative pressure from vacuum line 3 to the suction ports 2 contained within the block. The negative pressure is preferably imposed by one or more vacuum lines 3 which may run parallel to the shaft 21 before terminating in the block 23 containing the suction ports 2 or at any convenient point in the suction port assembly 22. In a preferred embodiment, the blocks 23 are substantially rectangular structures wherein the suction ports 2 are placed in an array as described below.

Referring to Figure 6, the block 23 is comprised of a plurality of suction ports 2 which are connected to vacuum line 3 to create a negative pressure at each suction port 2. The number of individual vacuum lines 3 may be any number less than or equal to the number of suction ports 2. An individual suction port 2 may be connected to negative pressure via the pressure conducting space 4 (See Figure 7a) or by a passage 26 which communicates an individual suction port 2 directly to the vacuum line 3. The passage 26 is preferably smaller in circumference than the opening of the suction port 2. In the embodiment of Figure 6, every suction port 2 is connected to a vacuum line 3 which traverses substantially the length of the block 23. Alternatively, every other suction port may also be connected to either of two vacuum lines as shown in Figure 8. In the embodiment of Figure 8, the passages 26 are oriented off-center to the circumference of the opening of the suction port 2.

Referring to Figure 7, a sectional view of the block 23 through line B-B of Figure 6 shows the suction ports 2 disposed within the block 23 in a linear array such that

tion port 2. The openings 12 of the suction ports 2 are flush with the bottom surface 6 of the block 23. In this embodiment, a pair of vacuum lines 3 connects each suction port 2 to negative pressure via a single passage 26 which runs from the vacuum line 3 to the suction port 2 and is substantially perpendicular to the vacuum line 3. Alternatively, as shown in Figure 8, vacuum lines 3 may connect each suction port 2 to negative pressure.

Referring to Figure 7a, the block 23 may also have an inlet 5 which introduces the negative pressure to the suction ports 2. The suction ports 2 are disposed within the bottom surface 6 of the block 23 which is to be affixed to the end of the shaft 21 (see Figures 5 and 9). As will be readily appreciated, the negative pressure can be introduced to the suction ports 2 by a variety of techniques as described above. The vacuum line 3 may be attached to the apparatus 20 at several points such as the individual suction ports as illustrated in Figure 7 or via a single inlet 5 and communicated to each suction port 2 via pressure conducting space 4 (Figure 7a) such that the negative pressure is introduced to the block 23 and communicated to each suction port 2. Thus, the objective of communicating the negative pressure to the suction ports 2 to fix the position of the cardiac tissue may be achieved via several alternate techniques which are based on or may be developed from the embodiments disclosed herein.

Referring to Figures 9 and 9a, as noted above the block 23 may be removable from the apparatus 20. The terminal and distal end of the apparatus 20 has a receiving means 29 which is adapted to reversibly receive the suction port assembly and the block 23. In a preferred embodiment, the block 23 is a separate molded rubber unit which is configured to fit reversibly into a receiving means 29 at the terminal end of the shaft 21. In this configuration, the suction port assembly 22 may be disassembled and the block 23 disposed after a single use. The receiving means 29 is comprised of a substantially parallel walls 27 having a grip means 28 associated therewith to maintain the position of the block 23 in place when attached to the receiving means 29. The grip means 28 may be one of several alternate designs. For example, the block 23 may have a ridge or indentation 30 which runs along the lateral exterior surface of the block 23 and fits conformingly in a groove or ridge 31 formed in the interior face of the wall 27.

The particular examples set forth herein are instructional and should not be interpreted as limitations on the applications to which those of ordinary skill are able to apply this invention. Modifications and other uses are available to those skilled in the art which are encompassed within the spirit of the invention as defined by the scope of the following claims.

Claims:

1. A device for stabilizing the beating heart by imposing a negative pressure on the surface of the heart

shaped to engage the surface of the heart, a plurality of suction ports having openings disposed in the bottom surface of the housing, and a vacuum line operably connected to the suction ports.

2. The device of claim 1 further comprising a means for introducing a negative pressure located in the interior of the housing.
3. The device of claim 2 wherein the means for introducing a negative pressure is comprised of a pressure conducting space connected to an inlet and to each suction port.
4. The device of claim 2 wherein the housing is annular and has a plurality of suction ports disposed about the periphery.
5. The device of claim 4 further comprising at least one instrument port located in the annular housing.
6. The device of claim 2 wherein the housing is dome-shaped.
7. The device of claim 6 wherein the means for introducing a negative pressure is comprised of an inlet and a pressure conducting chamber connected to each suction port.
8. The device of claim 6 further comprising at least one instrument port located in the dome-shaped portion of the housing.
9. The device of claim 8 wherein the instrument port is further comprised of means for attaching an instrument.
10. A device for stabilizing the beating heart by imposing a negative pressure on the surface of the heart comprising:
a shaft attached to a suction port assembly and means for introducing a negative pressure to the surface of the beating heart.
11. The device of claim 10 wherein said suction port assembly is comprised of a block having a plurality of suction ports disposed therein, and wherein said block is adapted to receive a vacuum line.
12. The device of claim 11 wherein the openings of the plurality of suction ports are disposed in the bottom surface of the block.
13. The device of claim 11 further comprising means for introducing a negative pressure contained within said block and connected to each of said plurality of suction ports.

14. The device of claim 13 wherein the means for introducing a negative pressure is comprised of a pressure conducting space pneumatically connected to said plurality of suction ports.

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15. The device of claim 10 comprising first and second shafts wherein each of said first and second shafts are attached to a suction port assembly.

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16. The device of claim 11 wherein each of said plurality of suction ports of said suction port assembly is connected to said vacuum line by a pressure conducting space in pneumatic communication with the interior of said block.

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17. The device of claim 16 wherein the pressure conducting space has an inlet attached to a vacuum line.

18. The device of claim 15 wherein said first and second shafts are interconnected and have proximal grips adapted to be held by hand.

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19. The device of claim 10 wherein said shaft has a receiving means at the distal end thereof for releasable attachment of said suction port assembly.

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20. The device of claim 19 further comprising said suction port assembly attached to said receiving means.

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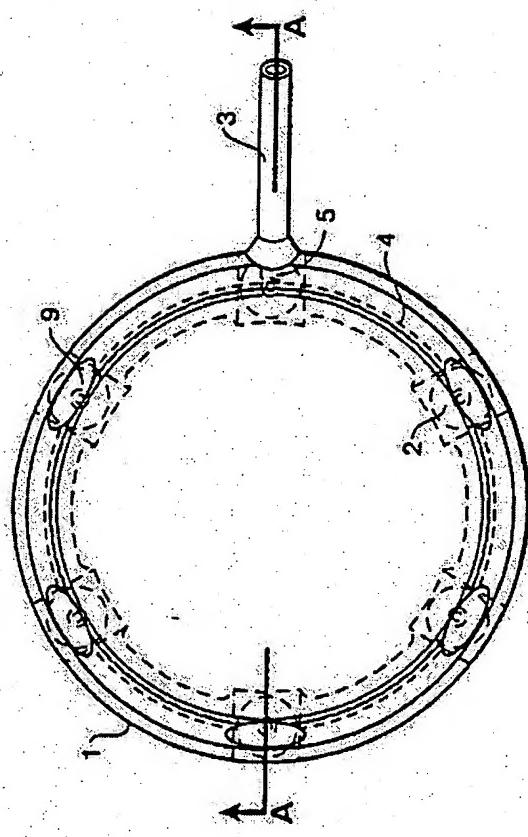


FIG. 1

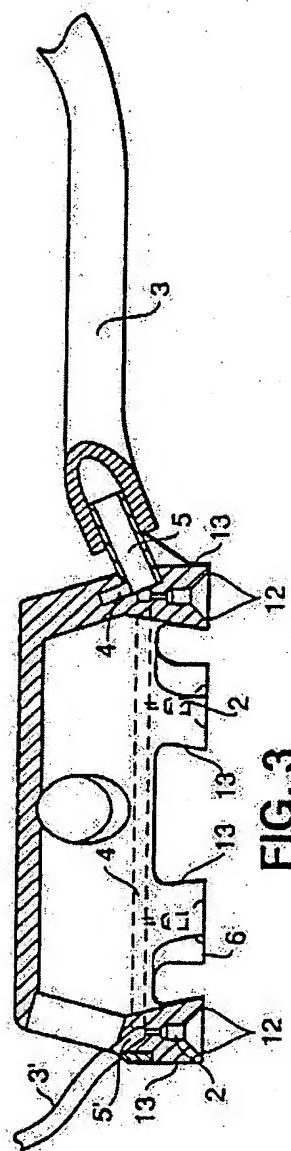


FIG. 3

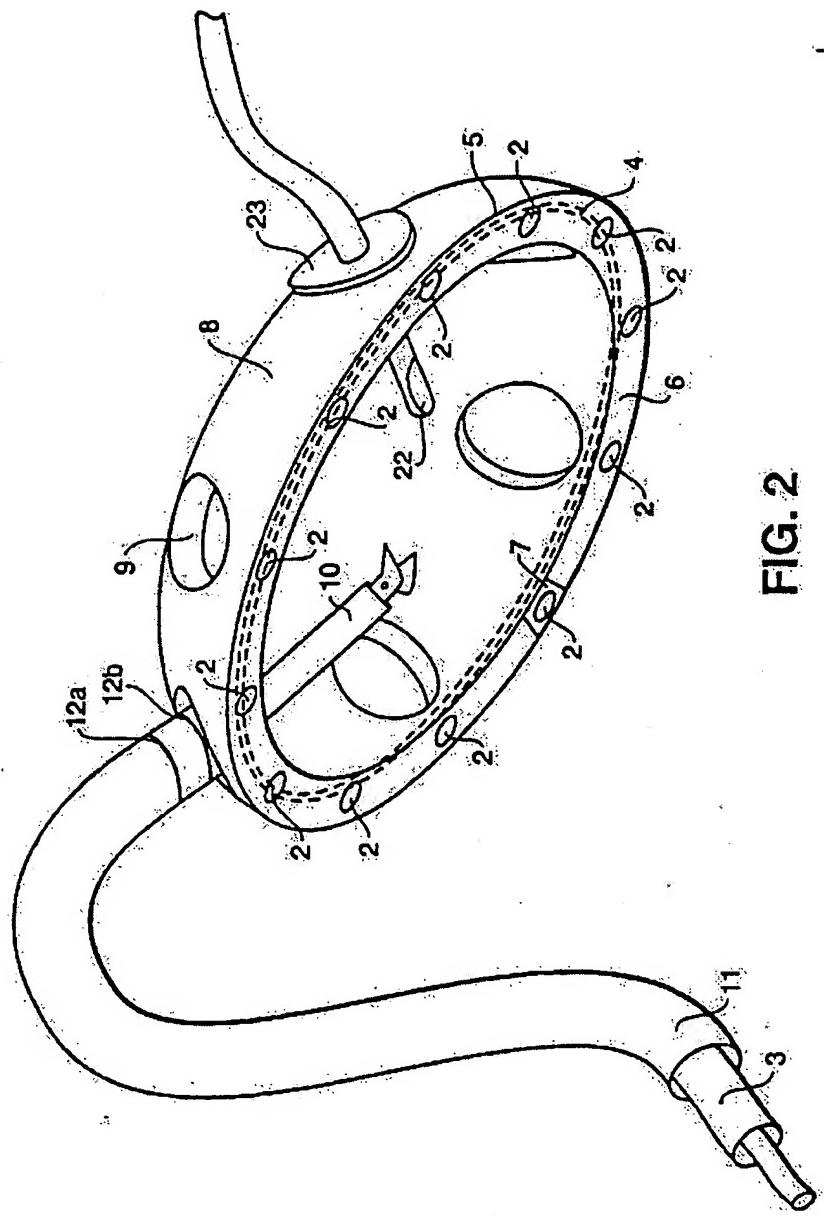


FIG. 2

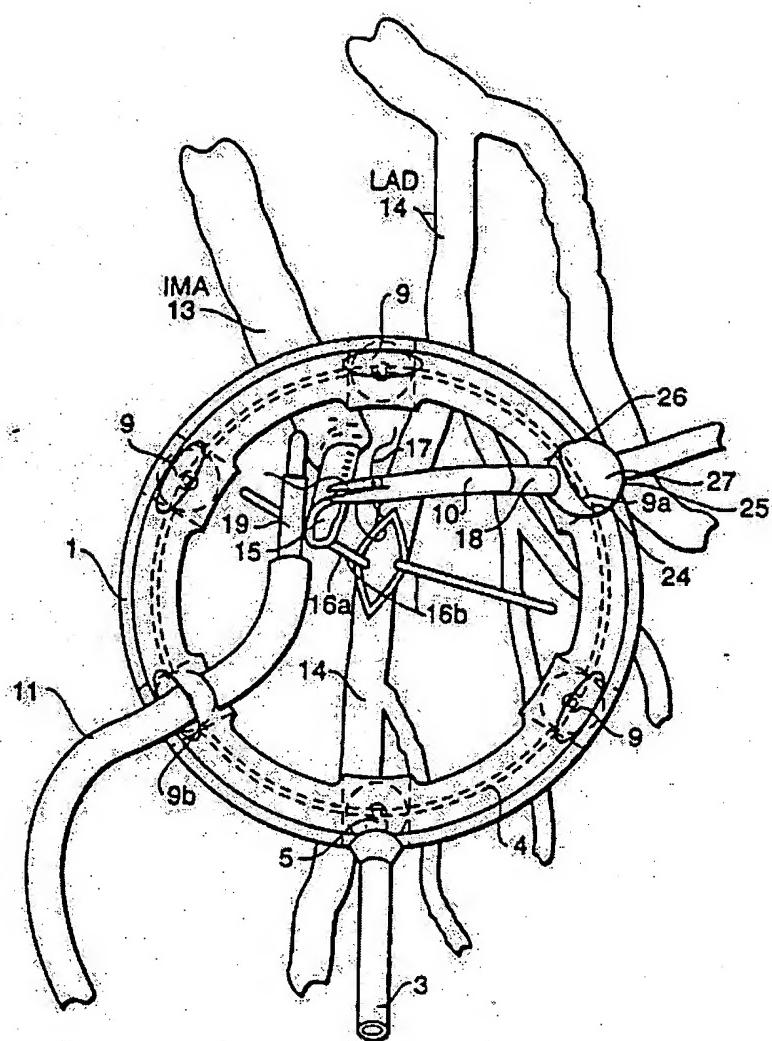


FIG. 4

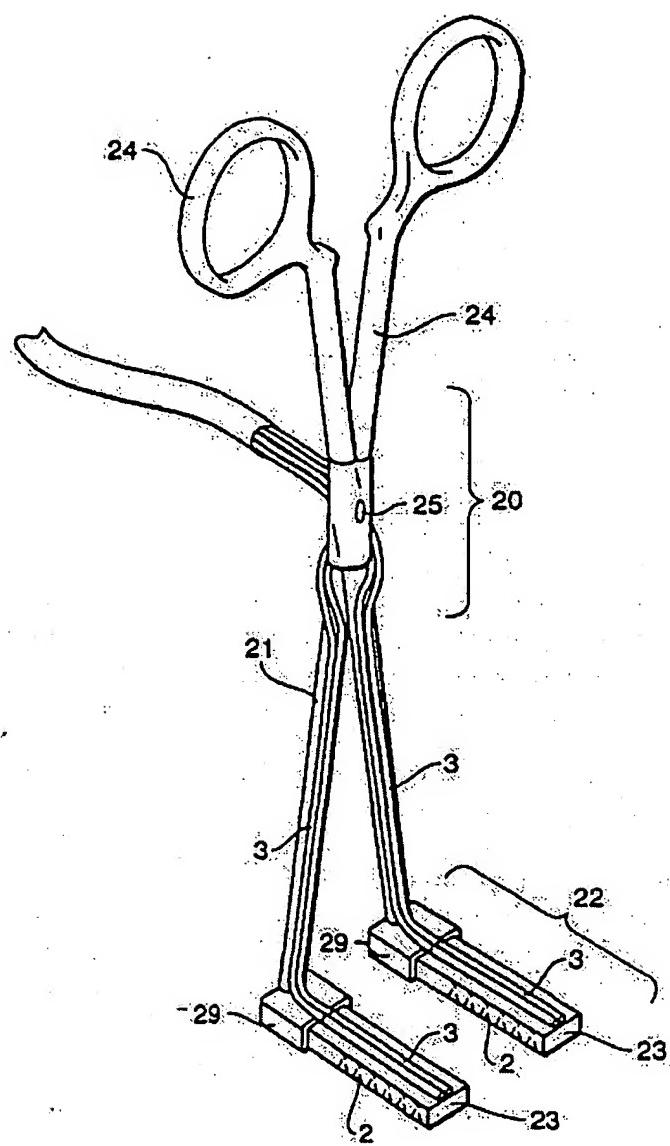
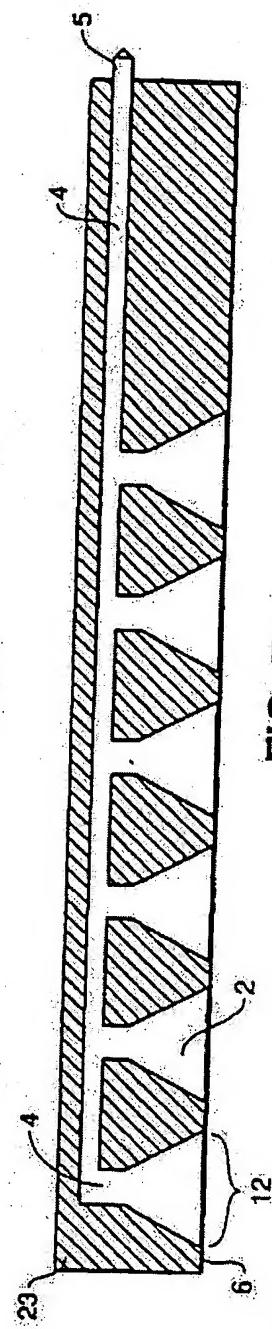
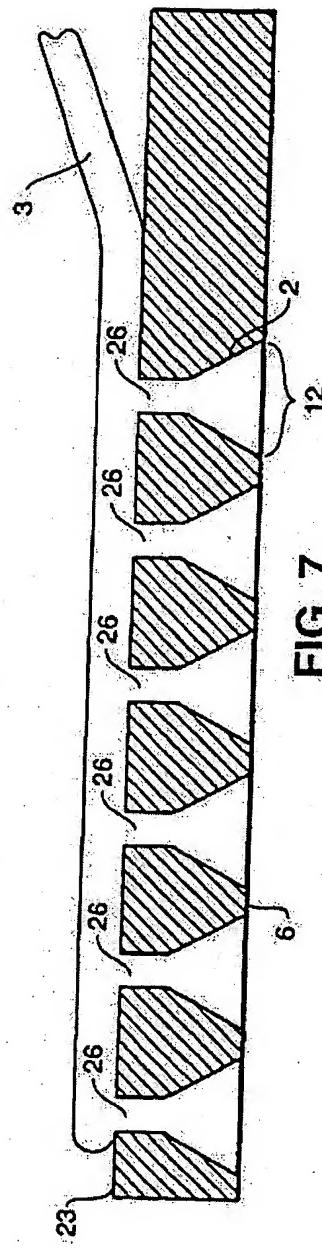
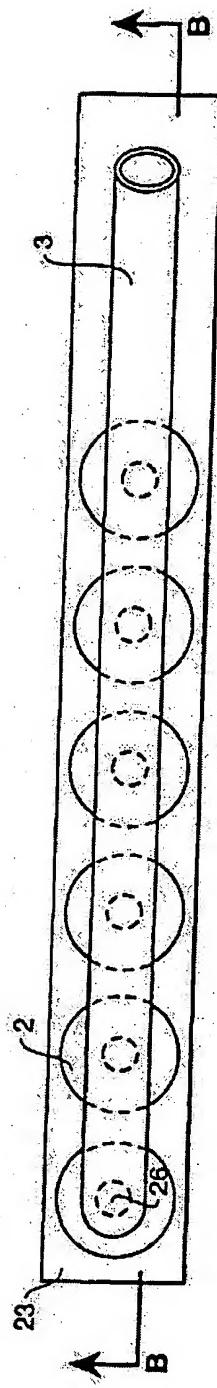


FIG. 5



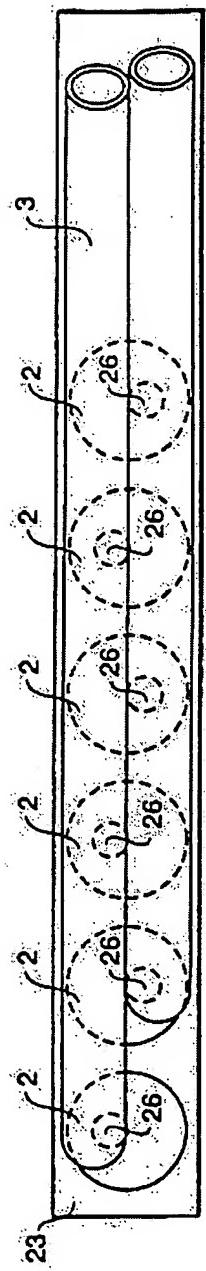


FIG. 8

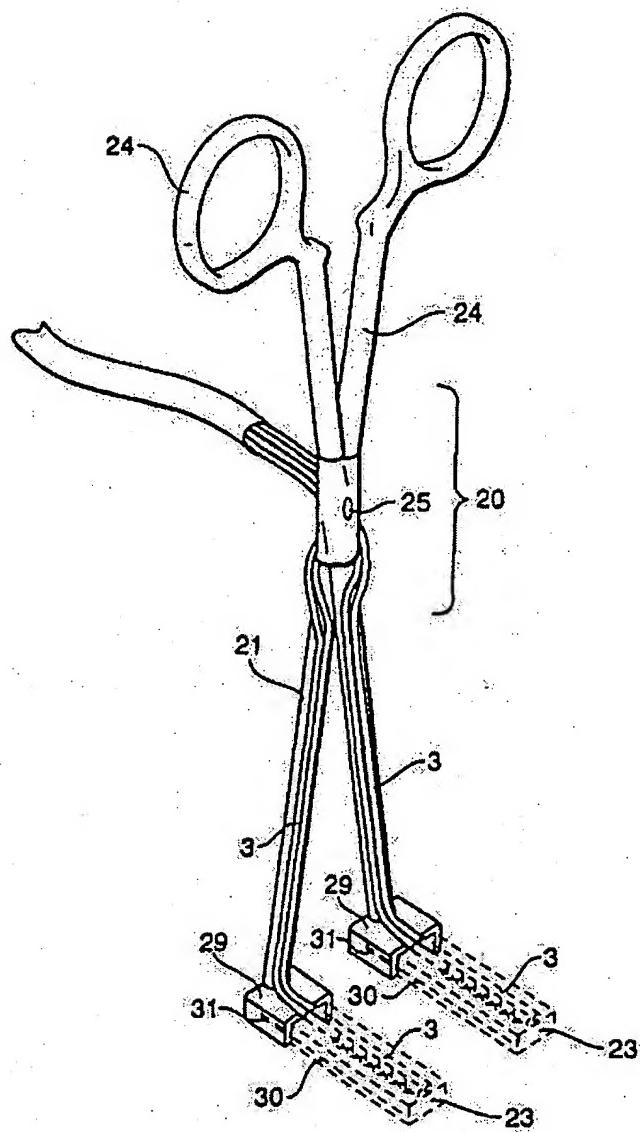


FIG. 9

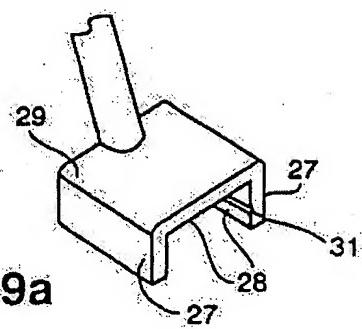


FIG. 9a



EUROPEAN SEARCH REPORT

Application Number

EP 97 10 2790

DOCUMENTS CONSIDERED TO BE RELEVANT									
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.)						
Y	WO 95 17127 A (OTICON AS ; RYGAARD JOERGEN A (DK)) 29 June 1995 * page 8, line 26 - page 9, line 6 *	1,10	A61B17/00						
A		2,5							
Y	DE 90 04 513 U (HERZBER WOLFGANG) 28 June 1990	1,10							
A	* page 5, line 6 - page 8, line 18 *	2-9, 11-20							
A	EP 0 668 058 A (NOVOSTE CORP) 23 August 1995 * column 8, line 37 - column 9, line 47 *	1,2,5,10							
A	WO 87 04081 A (LICENCIA TALALMANYOKAT) 16 July 1987 * page 4, line 21 - page 5, line 25 *	1,2,5,10							
TECHNICAL FIELDS SEARCHED. (Int.Cl.)									
A61B									
<p>The present search report has been drawn up for all claims</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 33%;">Place of search</td> <td style="width: 33%;">Date of completion of the search</td> <td style="width: 33%;">Examiner</td> </tr> <tr> <td>THE HAGUE</td> <td>9 May 1997</td> <td>Verelst, P</td> </tr> </table>				Place of search	Date of completion of the search	Examiner	THE HAGUE	9 May 1997	Verelst, P
Place of search	Date of completion of the search	Examiner							
THE HAGUE	9 May 1997	Verelst, P							
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document									
T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application I : document cited for other reasons & : member of the same patent family, corresponding document									

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